

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
21 October 2004 (21.10.2004)

PCT

(10) International Publication Number
WO 2004/089210 A1

(51) International Patent Classification⁷: **A61B 5/0452**

(21) International Application Number:
PCT/FI2004/050034

(22) International Filing Date: 2 April 2004 (02.04.2004)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
20030547 10 April 2003 (10.04.2003) FI

(71) Applicant and

(72) Inventor: KORHONEN, Pentti [FI/FI]; Vaahteratie 6,
FIN-36200 Kangasala (FI).

(74) Agent: TAMPEREEN PATENTTITOIMISTO OY;
Hermiankatu 12 B, FIN-33720 Tampere (FI).

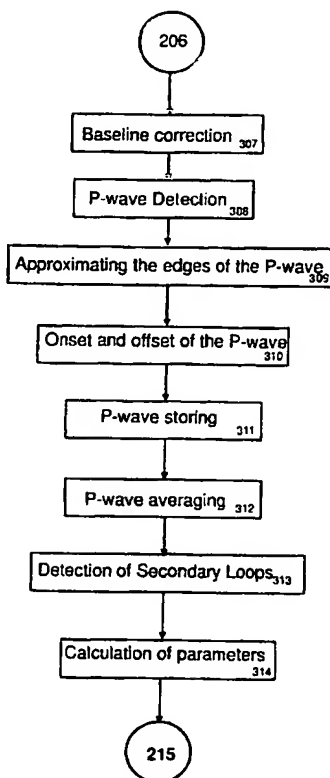
(81) Designated States (unless otherwise indicated, for every
kind of national protection available): AE, AG, AL, AM, — Published:
with international search report

AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN,
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,
GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE,
KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD,
MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG,
PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM,
TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM,
ZW.

(84) Designated States (unless otherwise indicated, for every
kind of regional protection available): ARIPO (BW, GH,
GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), Euro-
pean (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR,
GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK,
TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW,
ML, MR, NE, SN, TD, TG).

[Continued on next page]

(54) Title: SYSTEM AND METHOD FOR ANALYSING THE P-WAVE OF AN ECG-SIGNAL



(57) Abstract: The invention relates to a method, to a system and to a computer program product for the cardiac analysis. The method comprises steps for acquiring the ECG-signal in the form of vectorcardiography, detecting wave from the ECG-signal by a template method and calculating the parameter values of the P-wave preferably continuously during the ECG-recording. The method is aimed to the dynamic changes of the Rwave, wherein substantially every detected P wave is compared to the reference P-wave in defined time period. The cardiac analysis system according to the invention is configured to implement the aforementioned method. The computer program product comprises computer program code for implementing the aforementioned method.



— before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

6/PRTS

JC05 Rec'd PCT/PTO 07 OCT 2005

1

SYSTEM AND METHOD FOR ANALYSING THE P-WAVE OF AN ECG-SIGNAL

The invention relates to a method according to the preamble of the appended independent claim 1 for cardiac analysis, which method comprises steps for acquiring an ECG-signal, detecting at least one wave of the ECG-signal and calculating parameter values of said wave. The invention also relates to a cardiac analysis system according to the preamble of the appended independent claim 19 for implementing the aforementioned method as well as to a computer program product according to the preamble of the appended independent claim 32.

Background of the invention

An electrocardiogram (ECG) is a record of electrical events occurring within the heart of a patient and it is used for examination of heart diseases. The ECG plots the sequence of changes in electrical potential differences recorded between different regions of the patient's body surface. The ECG can be measured using various lead systems known as such. Commonly the ECG is obtained by using a standard 12-lead arrangement, but it can be obtained by using other lead systems, for example a Frank system, which uses three leads. These ECG-measurements provide a one-dimensional electrocardiographic signal, but it is also possible to present ECG as a three-dimensional signal, a so-called vectorcardiography.

The basic form of the ECG-signal can be seen in Figure 1. The ECG breaks down each heartbeat into a series of electrical waves. Three of the waves (P, QRS, T in the figure), a P-wave, a QRS-complex and a T-wave, are associated with the heart's contractions. The first deflection of the ECG is called the P-wave. It reflects the sequential activation (depolarization) of the heart's upper chambers, the right and the left atria. After the P-wave the ECG returns to its baseline, which is a substantially straight line on an ECG-paper where there are no positive or negative changes of electricity to create deflections. The next is a complex called QRS-complex, which reflects the sequential activation of the heart's lower chambers, the right and the left ventricle.

After the QRS-complex, the ECG returns to, or very nearly to, its baseline which is called a ST-segment, where it remains until the appearance of a T-wave. The T-wave reflects ventricular repolarization.

- 5 When discussing the ECG, a term "interval" refers to the length of the wave along with an isoelectric line following it. The interval can be named by using the letters of both waves on either side (e.g. PQ). A space between dash lines 1 and 3 (Figure 1) represents the PQ-interval. A term "segment" refers to the baseline between the end of
10 one wave and the beginning of the next wave. The space between dash lines 2 and 3 represents the PQ-segment, as well as the space between dash lines 4 and 5 represent the ST-segment.

- The above-mentioned vectorcardiogram (VCG) is a form of the
15 electrocardiography that represents movements of vectors of the myocardial activation with direction and magnitude as loops in a three-dimensional space. Vectorcardiography can be considered as a method of recording the direction and magnitude of electrical forces of the heart by means of a continuous series of vectors that form a
20 curving line around a center. A spatial orientation and magnitude of the heart vector is projected onto three orthogonal planes as frontal, horizontal and sagittal planes. The tip of the vector with coordinates (X, Y, Z) should trace out a loop in space, called a vector loop. The vector loop starts from the zero-point, which corresponds to the isoelectric line
25 or the baseline on a usual scalar ECG. The loop consists of, similar to the ECG-signal, a P-loop, a QRS-loop and a T-loop. When heart cycle is completed, the QRS-loop returns, in a normal situation, to zero-point and closes the loop. On the other hand, if the patient has a ST-change, the QRS-complex will not end to the baseline, but over or under,
30 depending on whether the patient has a ST-elevation or depression.

- The general features of the QRS-complex and the ST-segment within the ECG-signal have been extensively studied and their diagnostic
35 significance is quite well known. On the contrary, far less attention has been paid to the P-wave and therefore the clinical properties of this waveform are still not fully understood. It is known that the P-wave

represents atrial depolarization and it can show the heart's rate and rhythm. It is known that the atrial abnormalities and the P-waves are related to some heart diseases, for example to heart failure or to the susceptibility to atrial arrhythmias. Still, the specificity of these changes has been low.

Developed ECG-systems observe the QRS-complex and the ST-segment ideally, but have a lack in their operation of observing the P-wave. A reason for that can be found, in common opinion, where the P-wave and the atrias have been considered less meaningful compared to the QRS-wave, the ST-segment and the ventricles. Due to this, there are significantly less studies concerning the P-wave. In addition, the known studies of the P-wave use mostly the known 12-lead ECG. Such a 12-lead ECG-arrangement has a disadvantage in expression of the waves; it does not give a good three-dimensional picture of the electrical waves, which neglects the small waves, for example the P-wave. Studies done of the P-wave utilize mostly the ECG recording system that expresses the heart situation in one moment, but is not capable of continuously showing the in-time coming dynamic changes of the examined P-wave.

Short description of the invention

The present invention relates to a cardiac analysis method and system, which takes into account said lacks in existing systems. More precisely, the cardiac analysis method according to the invention comprises the steps for acquiring the ECG-signal, detecting at least one wave of the ECG-signal and calculating the parameter values of said wave, which is the P-wave. The cardiac analysis method according to the invention is then focused to the dynamic changes of the P-wave. The cardiac analysis system according to the invention comprises means for acquiring the ECG-signal, means for detecting at least one wave of the ECG-signal and means for calculating the parameter values of said wave, which is the P-wave. The system is additionally adapted to focus the analysis to the dynamic changes of the P-wave, wherein said system also comprises means for comparing substantially every

detected P-wave to the reference P-wave in defined time period. The computer program product according to the invention comprises computer readable instructions for detecting at least one wave from the ECG-signal, which said wave is a P-wave, whereupon the computer
5 program code comprises computer readable instructions for focusing to the dynamic changes of the P-wave, wherein said computer program code additionally comprises computer instructions configured to compare substantially every detected P-wave to a reference P-wave in defined time period.

10. In accordance with the invention, the ECG-signal is processed in the form of vectorcardiogram that can be described by three orthogonal leads: X, Y, Z. P-wave is at first recognized by a template and then the signals of the P-wave are averaged to form a smooth wave form, which
15 is used to determine the P-wave specific parameter values continuously from the whole ECG-recording. The cardiac analysis method according to the invention is then addressed to the dynamic changes of said P-wave. That means that it addresses to the changes of P-wave configuration that happen smoothly in the follow-up time.
20 The method comprises means for comparing substantially every detected P-wave to the reference P-wave in a defined time period. The results from each parameter are presented as points on a trend graph. The cardiac analysis method according the invention also provides means to notice and pick up the atrial extrasystoles, to separate them
25 into different sub-classes based on their morphology and then to average and analyze them in the same way as the P-wave.

The analysis of the P-wave according to the invention, brings new tools to the analysis of atrial activation; in consequence the P-wave is
30 considered to be more meaningful than before. With the method according to the invention, there is a first-time chance to see the dynamic changes of P-wave in a patient who has an acute myocardial infarction that affects also the atrial tissue. The atrial infarction is the entity that is supposed to affect the patient outcome by causing atrial
35 arrhythmias and ruptures of the atrial free wall in special situations. Still,

the dynamic changes in the P-wave configuration and the diagnostic criteries of the atrial infarction are unclear.

5 The analysis of the P-wave according to the invention provides also an effective way to monitor atrial arrhythmias, especially the paroxysmal atrial fibrillation. The atrial fibrillation is the most common cardiac arrhythmia, the prevalence of which is increasing with the aging of the population. Because of its clinical importance and the lack of highly
10 satisfactory management approaches, it is the subject of active clinical and research efforts. With the method according to the invention, it is possible to observe the dynamic changes of P-wave after the cardioversion when the normal sinus rhythm has been achieved and to try to find explanations for why some of the patients will maintain sinus
15 rhythm after the cardioversion and some of them will not. Additionally, with the method according to the invention, it is possible to observe the dynamic changes of P-wave in situations where the risk of atrial fibrillation is high, for example after heart surgery. Because the method of the invention is based on dynamic changes of the P-wave, it also gives a possibility of observing the actions of treatment and medicine
20 that is given to maintain sinus rhythm.

The analysis of the P-wave according to the invention also provides an effective way to monitor the dynamic changes of P-wave in the patient who has an acute heart insufficiency. These P-wave abnormalities are
25 discussed in more detail later in the description. It is obvious that with the method according to the invention, it is possible to observe the dynamic changes of P-wave also in other situations where the atrias are in the changing abnormal situation that affects the P-wave configuration as well.

30 The preferred embodiment of the invention is set forth in the drawings, in the detailed description which follows, and in the annexed claims. Further objects and advantages of the invention are also considered in the description. The invention itself is defined with particularity in the
35 claims.

Brief description of the drawings

- Figure 1 is a graphical representation illustrating the ECG-signal,
- 5 Figure 2 illustrates a flow chart of the steps of the method according to the invention,
- 10 Figure 3 illustrates a flow chart of the steps of the P-wave analysis according to the invention,
- Figure 4a-c illustrates the P-wave in one-dimensional environment,
- 15 Figure 5a-b illustrates the P-wave vector loop in two-dimensional environment,
- Figure 6a-b illustrates the P-wave vector loop in three-dimensional environment,
- 20 Figure 7 illustrates the P-wave in magnitude ECG, and
- Figure 8a-b illustrates the change area, the change vector angle and magnitude.

25

Detailed description of the preferred embodiment

- Figure 2 illustrates the steps of the method according to the invention as a flow chart. The present invention exploits the known three-dimensional electrical ECG-model called vectorcardiogram. ECG-signal acquisition (201) can be done in various ways. A basic method is to use eight standard ECG-surface electrodes, which are placed on the patient according to the Frank electrode system. The electrodes are then used to form an ECG-vectorcardiogram by method known in prior art, which ECG-vectorcardiogram can be described by three orthogonal leads X, Y, Z. It is also possible to acquire the ECG-signal by using a
- 30
- 35

standard twelve-lead-ECG-arrangement, which is stored and derived to form a vectorcardiogram and then further analyzed. According to the invention, the ECG-signal is acquired preferably from existing vectorcardiogram data collected by another system (200). For example, the ECG-signal can preferably be acquired from the MIDA data storage unit. The MIDA is a commercial monitoring system (sold by Philips Medical) made to analyze the ischemic changes of QRS-complex and ST-segment of ECG and it is widely used in hospitals. MIDA registers the electrical signals of the heart using said Frank electrode system and constructs the three-dimensional electrical model from them. The action of MIDA is discussed in more detail in US-5520191. It is obvious that the ECG-signal in the present invention can be acquired by not only the mentioned but any known methods.

After the acquisition, the ECG-signal is preprocessed (202) to minimize electrical artefacts. Preferably the raw data is filtered to remove the noise and to improve its signal-to-noise-ratio (SNR). The noise is filtered by some known signal processing method, which are not discussed more in this as it can be considered obvious to the man skilled in signal processing.

The preprocessed signal is according to the invention analyzed to detect the peaks of R-waves (203). A gap between two peaks is referred to by a term "beat". When the peaks are found, a time period between them is measured and the beat is stored. Beats are classified depending on their duration (time period). If the time between two consecutive Rwaves changes suddenly under predetermined time, it can be assumed that the beat consists of an atrial extra systole (204). In that case the beat is stored to the atrial extra systole database for further analysis (205). The atrial extra systoles can be categorized into sub-classes according to their morphology. Afterwards, they are averaged and analyzed. The result of this examination will give information of the changes of the atrial extra systoles as well as the number of the atrial extra systole sub-groups. By picking the extra systoles from the ECG-signal, the resulting R-wave data will become as homogenous as possible, which is preferable to the analysis. There

is also a possibility that the change results from the ventricular extra systole when there is no atrial extra systole preceding the deviant-looking QRS-complex. In this case, the changed beat is also picked to the second database and later removed. All the other beats (the unchanged, time period within the predetermined time limit) are stored to a P-wave database. The beats in said both databases are stored in the X, Y, Z -leads (206) and used separately for the P-wave analysis according to the invention.

10 P-wave analysis:

Reference is now made to Figure 3, in which the examination of the P-wave is described in more detail. At first the P-wave signal is processed by baseline correction (307) to minimize the baseline drift of the signal. This way the signal quality will improve. There are several methods for implementing the baseline correction, for example linear interpolation, quadratic or cubic corrections, polynomial fitting and high pass frequency filtering. Because said methods are known as such, they are not explained further.

20 The P-wave is normally viewed as a smooth, small and curved deflection. It is usually a positive deflection, although it may be negative as well. The duration of the P-wave is normally below 0.12 seconds and the amplitude is normally less than 0.25 mV. In the frontal plane the P-wave axis angle ranges from 0 to 75 degrees. Also notched P-waves may be seen and the normal P-wave may often exhibit two components as an M-shaped complex. Because of its small, ill-defined and variable shape, the detection of the P-wave is difficult.

30 According to the invention, the P-wave detection (308) is preferably done by a template method, which is a type of two-dimensional cross-correlation method with both time and amplitude as variables. The template method attempts to detect the P-wave by using the covariance between the template signal and the wave form signal stored beat. For substantially each beat, the template is tried to be

35

matched with the actual signal and the P-wave is detected when the covariance exceeds a specified threshold. The P-wave is valid if its duration is between the minimum and the maximum P-wave durations. As said, the length of the P-wave is generally about 120 ms. Time
5 between the beginning of the P-wave to the beginning of the Q-wave (called PQ-time) is generally below 200 ms. These values are preferably used to form a template time window, but the time window can be predefined with some other values or with entering the values occasionally.

10 The P-wave detection is preferably carried out only in one coordinate, because frequently one coordinate is less noisy than the others and therefore more suitable for analysis. In the detection, the user is at first asked to enter the starting and ending sample of the template in order
15 to construct the first template. Later, the template can be changed if there is a need for it. It can be done manually in the same way as the first template. It can be done also automatically in that way that the predetermined amount of latest detected P-waves is taken and averaged, wherein the averaged P-wave is set to be the next template.
20 The template is also checked for the adaptation, if the amount of dropped P-waves exceed the predetermined level. In these ways, the template can be changed according to the changes of the P-wave. In other words, template changes as the P-wave changes. Obviously, the P-wave can also be detected by some other methods known in the
25 prior art, such as thresholding, pattern recognition etc.

It has been empirically observed that the preprocessed ECG-signal can still be relatively noisy around the P-wave, which can affect the determination of the real location of the onset and offset of P-wave.
30 The amplitude of the deflection might be affected by the existence of a noise plateau before the beginning of the P-wave. For avoiding this an improved method of detection of the P-wave time limit is preferably used (309). In the improved method, the rising and falling edges of the P-wave are approximated by straight lines using a slope function. Later
35 values of the baseline of the beat are calculated and stored.

Intersection points of the baseline and the two edges are obtained by an intersection function.

5 With the help of the baseline, the value of the onset and the offset will be determined (310). The onset is the intersection of the baseline and the rising edge, whereas the value of the offset will be the intersection of the baseline and the falling edge. The value for the P-wave offset can also be found by a threshold method, in which the sample having the ECG of approximately the same value as in the P-wave onset is
10 searched. The found value is accepted as the real offset of the signal based on the idea that in most of the cases the P-wave bop will be closed in three-dimensional environment, which means (in time domain) that the P-wave must return to its initial value.

15 The method of finding the P-wave onset and offset is extremely important in analyzing the P-wave specific parameters, because in those calculations the P-wave must have clear boundaries, in other words, a P-wave vector loop must be completed. The P-wave vector loop is discussed in more detail later in the description. However, it is
20 equally important to notice that the offset function is not used while analyzing the PQ-segment specific parameters, because with those parameters the PQ-segment elevation, in other words the incomplete P-wave vector loop, must be analyzed in the unedited situation. Therefore, the P-wave onset function is used in the method
25 substantially all the time, and the P-wave offset function is used with the P-wave specific parameters.

The detected P-wave is stored in X-, Y- and Z-leads (Figure 4a, P-wave circled; Figure 3, 311). All the P-wave beats are then averaged
30 (312) in the predetermined time intervals to form a smooth beat. Formed beats are used for calculations and analysis. The first averaged P-wave is used as an initial reference P-wave, where the upcoming averaged P-wave is compared to (monitoring of the dynamic changes).

35

When monitoring the myocardial infarct, the patient is in cardiac monitor, e.g. MIDA, preferably for 1-2 days. Substantially all the ECG-signals during that time should be taken for the analysis. Averaging for the data is preferably done every 4 minutes.

5

When monitoring the cardioversion of the patient with atrial fibrillation, the normal sinus rhythm is observed preferably for 3-4 hours. Averaging for the data acquired is preferably done every 10 sec – 2 minutes. The reason for the shorter averaging time is due to the lesser data and the quicker changes in the beginning.

10

It is also possible to monitor the P-wave changes, for example, 1, 3 or 6 months after the procedure such as, for example, the cardioversion. In that case the monitoring may last 15-30 minutes and it focuses on observing the change of the vector loop. The different monitoring results are combined for better analysis.

15

It is obvious that the monitoring depends on the situation, when also the above-mentioned time limits vary according to the situation.

20

In some situations, the P-wave can be seen as two loops, the primary (A_p) and the secondary (A_{ip}) loop, which are detected (313). The loops are illustrated in figures 4b and 5b in one-dimensional and two-dimensional spaces. Detection of the secondary loop is essential in P-wave loop analysis, as it can be greatly valuable in the analysis of atrial anatomic changes described later in the text. P-wave loops in a three-dimensional space can preferably be projected onto the three orthogonal planes and the existence of the secondary loop can thus be completed in two dimensions.

25

30

The QRS-complex is also observed in the method according to the invention. This is done by detecting the beginning and the ending of the QRS-complex from the beginning of the beat. The duration of the QRS-complex is also measured, as well as the waveform signal. These values are used along the P-wave analysis.

35

According to the invention, the averaged P-waves are calculated and the parameters described later in the text are estimated next. (314). A few of the parameters are common parameters, several are developed for the invention and for the dynamic analysis of the P-wave, since they have not existed before. Parameters can be divided into four categories: one-dimensional environment, two-dimensional environment, three-dimensional environment and magnitude ECG environment.

10 One-dimensional environment

The parameters of a one-dimensional diagram describe the P-wave properties in one dimension and relate to the orthogonal axes of the electrocardiogram. Figures 4a and 4b illustrate the P-wave in a one-dimensional environment. A vector area (P-A) (Fig. 4a) expresses the area of the P-wave. It can be derived from the areas in the three orthogonal leads X, Y, Z to form an equation (sqrt corresponds to the square root):

$$20 \quad P-A = \sqrt{A_x^2 + A_y^2 + A_z^2}.$$

A vector change area (PC-A) describes the changes of the P-wave area. Preferably the area of the P-wave being examined (A_{exam}) is compared with the area of the reference P-wave (A_{ref}). The difference is calculated from X, Y, Z -leads:

$$PC-A = \sqrt{[(A_{exam} - A_{ref})_X]^2 + [(A_{exam} - A_{ref})_Y]^2 + [(A_{exam} - A_{ref})_Z]^2}.$$

A P-area duplicity (P-AD) can be calculated by relating the area of the secondary loop of the P-wave (A_{il}) to the primary loop area (A_j) from X, Y, Z-leads (Figure 4a).

$$P-AD = \sqrt{[(A_{il} / A_j)_X]^2 + [(A_{il} / A_j)_Y]^2 + [(A_{il} / A_j)_Z]^2}.$$

35 In addition, the PQ-parameters are calculated. A PQ-vector magnitude (PQ-VM) describes the PQ-magnitude (level of PQ segment elevation)

and a PQ-area (PQ-A) describes the area between the PQ-segment and the baseline (e) in a one-dimensional environment (Figure 4c). A PQC-A-parameter describes correspondingly the upcoming changes of the PQ-area compared to the values of the reference P-wave. A PQ-time and a P-wave duration (P-dur) are also calculated in the one-dimensional environment.

$$\begin{aligned} \text{PQ-VM} &= \sqrt{\text{PQ}_x^2 + \text{PQ}_y^2 + \text{PQ}_z^2} \\ \text{PQ-A} &= \sqrt{\text{PQA}_x^2 + \text{PQA}_y^2 + \text{PQA}_z^2} \\ 10 \quad \text{PQC-A} &= \sqrt{[(A_{\text{exam}} - A_{\text{ref}})_x]^2 + [(A_{\text{exam}} - A_{\text{ref}})_y]^2 + [(A_{\text{exam}} - A_{\text{ref}})_z]^2} \end{aligned}$$

Two-dimensional environment:

- 15 The parameters of a two-dimensional diagram describe a P-wave vector loop in the three orthogonal planes: frontal (XY), horizontal (XZ) and sagittal (YZ) planes. The P-wave vector loop in a two-dimensional environment is shown in figure 5a and 5b. A P-wave vector loop area (P-LA) is the area of a two-dimensional loop described by the P-wave vector (Figure 5a). It can be derived from the different areas in the
- 20 three orthogonal planes XY, XZ and YZ to form an equation:

$$\text{P-LA} = \sqrt{A_{xy}^2 + A_{xz}^2 + A_{yz}^2}$$

- 25 A P-wave change loop area (PC-LA) is the difference of the two-dimensional areas of the P-wave vector loop being examined and the reference P-wave vector loop:

$$30 \quad \text{PC-LA} = \sqrt{[(A_{\text{exam}} - A_{\text{ref}})_{xy}]^2 + [(A_{\text{exam}} - A_{\text{ref}})_{xz}]^2 + [(A_{\text{exam}} - A_{\text{ref}})_{yz}]^2}$$

A vector loop area duplicity (P-LAD) describes the P-wave duplicity ratio, which is the ratio of the secondary loop (A_{ij}) to the primary loop (A_i) (Figure 5b):

$$35 \quad \text{P-LAD} = \sqrt{(A_{ii} / A_i)_{xy}^2 + (A_{ii} / A_i)_{xz}^2 + (A_{ii} / A_i)_{yz}^2}$$

Three-dimensional environment:

The parameters of a three-dimensional diagram describe the P-wave vector loop in the three-dimensional environment (Figure 6a). A three-dimensional P-wave vector loop area (P3-LA) is the area of the three-dimensional loop described by the P-wave vector. The basic function, called "looparea", calculates the area of a loop dividing it into small triangles in three dimensions and using the known vector cross product method to compute their areas. The three-dimensional P-change loop area (P3C-LA) is the difference between the area of the P-wave vector loop being examined (A_{exam}) and the reference P-wave vector loop (A_{ref}):

$$P3C-LA = \sqrt{(A_{exam} - A_{ref})^2}.$$

A P-azimut (P-Az) is a parameter for the angle that a P-main vector (M) describes in the transversal plane (Figure 6b). The P-main vector is defined as the average vector of substantially all the P-wave vectors that compose a P-wave loop in three dimensions. A P-Elevation (P-El), shown also in Figure 6b, is a parameter for the angle that the P-main vector forms in the vertical plane.

A P-change vector angle (PC-VA) denotes the angle difference between the main vector of the P-wave examined and the reference P-wave. A P-VM-parameter is the P-vector magnitude and a P-change vector magnitude (PC-VM) is the magnitude of the vector point of the reference P-main vector to the examined P-main vector. A P-QRS-vector angle (PQRS-VA) expresses the angle between the P and the QRS-main vectors. A P-QRS-change vector angle (PQRSC-VA) expresses the upcoming changes compared to the values of the reference P- and QRS-waves.

A PQ-vector magnitude (PQ3-VM) describes the PQ-magnitude in a three-dimensional environment. A PQ-change vector magnitude (PQ3C-VM) is the distance of the vector point of the reference P-main

15

vector to the examined P-main vector magnitude in a three-dimensional environment.

- 5 In addition, angles of the PQ- vector (PQ-Az, PQ-EI) and the difference (PQC-VA) between the PQ-wave examined and the main vector of the reference PQ-wave are calculated.

- 10 A P-vector loop length (P-VLL) is the perimeter of the loop drawn by the P-wave vector in a three-dimensional space. A P-vector loop velocity (P-VLV) describes the speed of the development of the P-wave vector loop length.

Magnitude ECG:

- 15 The parameters of the magnitude ECG are a P-vector magnitude area and the difference of it. Figure 7 illustrates the P-wave in magnitude ECG. The P-vector magnitude area (P-MA) is an area of the P-wave in the magnitude ECG-signal. A P-vector magnitude area difference (PC-MA) describes the change between the examined P-wave magnitude area and the reference P-wave magnitude area:
- 20

$$PC-MA = \sqrt{(A_{exam} - A_{ref})^2}$$

- 25 In addition, the PQ-parameters are calculated similar to the one-dimensional environment. A PQ-MVM describes the PQ-magnitude and a PQ-MA describes the PQ-area of the PQ-elevation. PQC-MA describes correspondingly the upcoming changes of the PQ-area compared to the values of the reference P-wave.

30 $PQC-MA = \sqrt{(A_{exam} - A_{ref})^2}$

Figure 8a represents the change of the P-wave area (PC-A) and figure 8b represents the change vector angle (PC-VA), as well as the change vector magnitude (PC-VM).

35

Parameters described above are preferably also used with the analysis of the beat stored into the atrium extra systole database.

5 When parameters have been calculated, the results are displayed (215). According to the invention, the results are displayed from every averaged time interval as a new point in trend curve over time. An advantage of this presentation is that the results are easier to see and a conclusion is easier to draw from the changes of the P-wave.

10 Analysis of the parameters:

Parameters that concern the PQ-segment can have a great impact when studying the development of an acute myocardial infarction of atria. It is known that the manifestations of atrial infarction may include
15 elevation or depression of the Ta-segment (Ta represents atrial repolarisation like the ST-segment and T-wave represent ventricular repolarisation). The Ta-segment is usually obscured by the QRS-complex and the early part of the ST-segment, but it's abnormalities may affect the PQ-segment. The parameters PQ-VM, PQ-A, PQC-A, PQ3-VM, PQ3C-VM, PQ-MVM, PQ-MA, PQC-MA, PQ-Az, PQ-EI and
20 PQC-VA are developed to analyze these changes.

Parameters that concern the P-wave primary (A_I) and secondary (A_{II}) loop are developed most of all to evaluate the P-wave changes in heart
25 insufficiency. It is known that in heart insufficiency the P-wave may become peaked or notched, depending on the etiology of disease. Although the specificity of these changes varies, it is a common opinion that in the left atrial enlargement due to heart insufficiency the P-wave became notched and has a negative terminal part, so that the P-wave
30 has the appearance of the "fallen S". The size of a negative terminal part may correlate to the level of the left atrial enlargement and heart insufficiency. The parameters P-AD and P-LAD are developed to analyze these dynamic changes.

35 Parameters that concern the QRS-complex vector angle and its relation to the P-wave vector angle are developed to discover if the

change of P-wave vector angle is due to change of the heart position. The parameters PQRS-VA and PQRSC-VA are developed to analyze these changes.

- 5 Most of the parameters that concern the P-wave and PQ-segment area, the vector magnitude and angle, the PQ-time and the P-duration, the length and the velocity of the vector loop describe the character of P-wave vector loop widely. Their changes reflect the condition of the P-wave dynamically. For example, in the acute myocardial infarction of
10 atria these parameters will change due to the tissue damage of the atrium.

- There is some evidence that in the 12-lead ECG the P-wave duration and dispersion (difference between the longest and shortest P-wave
15 duration) may be greater in persons who are in a higher risk to develop atrium fibrillation. Although the specificity of these changes varies, there is some evidence that when these changes decrease with the medication, the risk also decreases. As these changes in a 3-dimensional environment can be seen as increased irregularity of the
20 P-wave vector loop, the above-mentioned parameters describe these dynamic changes ideally. Parameters are suitable also in other situations where dynamic changes in intra-atrial pressure, volume or conduction alter the P-wave configuration slowly. In addition, these parameters are useful in investigation of the quality atrial extra
25 systoles.

System:

- 30 A cardiac analysis system according to the invention comprises means for signal processing and counting parameters. The system is adapted to take raw ECG-signal from existent datasystem and to store it into a file. Said cardiac analysis system is arranged to convert the sample file into a binary file containing X, Y, Z -samples. The samples in the generated file are preferably 16 bits signed integers. Samples are
35 advantageously ordered in $X(i)$, $Y(i)$, $Z(i)$, $X(i+1)$, $Y(i+1)$, $Z(i+1)$, wherein

X, Y, Z refers to the three orthogonal components of the VCG-signal and "i" refers to the beat number.

5 An input-file also comprises information about the resolution of the system, samples taken per second, data type (preferably signed 16 bit integer), beginning of the time frame to be analyzed related to the whole duration of the file, end of the time frame to be analyzed and number of channels (preferably three, corresponding to the orthogonal axes X, Y, Z). Naturally, the input-file can include other information as
10 well.

The system also comprises means for preprocessing to remove the noise of the signal and to improve the SNR. Preprocessing is preferably done by filtering, which can be implemented with both low
15 and high frequency of the bandpass preprocessing filter. The system also comprises a structure, which focuses on the beat specifications. Preferably, specification is done by measuring time and duration of the peaks and the waves. It can be adapted to measure the previous R peak, the posterior R-peak, as well as the beginning of the range to
20 calculate the baseline. Also, it can be stored with information of the channel used for Rpeak detector and baseline correction, the rate to calculate the threshold for Rpeak detection, and the duration of every beat in samples.

25 The detected beat is stored into the system. Information of the beat, such as a sample number when the "i"-th beat starts in the entire input file; a sample number of the ending of the "i"-th beat in the input file; duration of the beat in samples; wave form signal of the stored beat; baseline value of the "i"-th beat; duration of the beat in seconds;
30 location of the R-wave peak in samples; validity of the P-wave (0 or 1) determined preferably by duration; P-wave onset; P-wave offset and length of the P-wave in samples are preferably also stored into the system.

35 A P-wave detector of the system is capable of validating the P-wave by knowing the threshold of covariance of the template method and the

minimum and the maximum P-wave durations in samples. The validity of the P-wave is determined by measuring P-wave duration, which should settle between the minimum and the maximum values.

- 5 The system is preferably implemented as hardware and software. Therefore the system also comprises a computer program for implementing the method according to the invention. The computer program comprises computer readable code for acquiring the ECG-data, detecting the P-wave from said data and analyzing said P-wave
10 according to the method of the invention.

- The method and the system describe the preferred embodiments of the P-wave analysis according to the invention. The main idea in the method is to analyze the dynamic changes of the P-wave in time.
15 Implementation of the system can be carried out in different ways. For example it will be appreciated that in some cases other ECG-model is utilized, e.g. ECG-data can be acquired from a monitoring system using 10-lead arrangement, and determines an ECG-model that corresponds to the 12-lead arrangement to which the analysis is
20 focused. Similarly, other parts of the ECG-model can be included to the analysis, e.g. the PQ-segment as described earlier.

By knowing this, it is obvious that the present invention is not limited to the description, but to claims discussed herein below.

CLAIMS:

1. A method for a cardiac analysis, the method comprising steps for acquiring an ECG-signal, detecting at least one wave of the ECG-signal and calculating parameter values of said wave, **characterized** in that, said wave is a P-wave, whereupon the cardiac analysis is focused to dynamic changes of the configuration of the P-wave, wherein substantially every detected P-wave is compared to a reference P-wave in defined time period.
2. The method according to claim 1, **characterized** in that, the cardiac analysis is focused also to dynamic changes of the PQ-segment.
3. The method according to claim 1 or 2, **characterized** in that the ECG-signal is in the form of a vectorcardiogram.
4. The method according to claim 1 or 2 or 3, **characterized** in that a beat between two R-peaks is examined, whereupon said beat is classified into groups depending on whether the beat is having a duration between the predetermined time limit or the beat is having a duration under the predetermined time limit, whereupon both said beats are analyzed separately.
5. The method according to one of the claims 1 – 4, **characterized** in that the P-wave is detected by a template method.
6. The method according to one of the claims 1 – 4, **characterized** in that the P-wave is detected by a pattern recognition method.
7. The method according to one of the claims 1 – 6, **characterized** in that the detected P-wave is stored in X, Y, Z leads.
8. The method according to one of the claims 1 – 7, **characterized** in that the detected P-wave is averaged in the predetermined time interval.

9. The method according to claim 8, **characterized** in that at least one averaged P-wave is used as an initial reference P-wave, where the upcoming averaged P-waves are compared to.
- 5 10. The method according to one of the claims 1 – 9, **characterized** in that at least one loop of the P-wave is detected.
- 10 11. The method according to one of the claims 1 – 10, **characterized** in that the parameters of the P-wave in one-dimensional diagram are one or more of the following: the vector area (P-A), vector change area (PC-A), P-area duplicity (P-AD), PQ-vector magnitude (PQ-VM), PQ-area (PQ-A) and PQ change area (PQC-A).
- 15 12. The method according to one of the claims 1 – 11, **characterized** in that the parameters of the P-wave in two-dimensional diagram are one or more of the following: the vector loop area (P-LA), vector change loop area (PC-LA) and P loop area duplicity (P-LAD).
- 20 13. The method according to one of the claims 1 – 12, **characterized** in that the parameters of the P-wave in three-dimensional diagram are one or more of the following: the vector loop area (P3-LA), the vector change loop area (P3C-LA), the angles of the azimuth (P-Az, PQ-Az), the elevation (P-EI, PQ-EI), change vector (PC-VA, PQRSC-VA, PQC-VA), the P-QRS vector (PQRS-VA) as well as the vector magnitude (P-VM, PQ3-VM), change vector magnitude (PC-VM, PQC3-VM).
- 25 14. The method according to one of the claims 1 – 13, **characterized** in that the parameters of the P-wave in magnitude environment are one or more of the following: the vector magnitude area (P-MA, PQ-MA), the vector change magnitude area difference (PC-MA, PQC-MA) and the vector magnitude (PQ-MVM).
- 30 15. The method according to one of the claims 1 – 14, **characterized** in that the method comprises also the calculations of one or more of the following: the PQ-time, P-wave duration (P-dur), the length of the P-wave (P-VLL), the velocity of the P-wave vector loop (P-VLV).
- 35

16. The method according to one of the claims 1 – 15, characterized in that the ECG-signal is acquired from a Frank system or a 12-lead ECG-arrangement.
- 5 17. The method according to one of the claims 1 – 16, characterized in that the ECG-signal is acquired from a data storage unit that is independent of this invention and commercially available.
- 10 18. The method according to one of the claims 1 – 17, characterized in that results of the parameters are displayed in a trend curve.
- 15 19. A cardiac analysis system comprising first means for acquiring the ECG-signal, second means for detecting at least one wave from the ECG-signal and third means for calculating parameter values of said wave, characterized in that, said wave is a P-wave, whereupon the cardiac analysis system is adapted to focus to dynamic changes of the configuration of the P-wave, wherein said system additionally comprises means for comparing substantially every detected P-wave to
- 20 a reference P-wave in defined time period.
20. The system according to claim 19, characterized in that, the system is further adapted to focus to dynamic changes of the PQ-segment.
- 25 21. The system according to claim 19 or 20, characterized in that the ECG-signal is in form of a vectorcardiogram.
- 30 22. The system according to claim 19 or 20 or 21, characterized in that the system is also adapted to measure a duration of the beat between two R-peaks, wherein the system is also configured to compare the beat to the predetermined time limit and classified the beat into the one of two groups depending on whether the duration is between the predetermined time limit or under the predetermined time limit, wherein the system is also configured to analyze both groups
- 35 separately.

23. The system according to one of the claims 19 – 22, characterized in that the system is adapted to detect the P-wave by a template method.
- 5 24. The system according to one of the claims 19 – 22, characterized in that the system is adapted to detect the P-wave by a pattern recognition method.
- 10 25. The system according to one of the claims 19 – 24, characterized in that the system is adapted to store the detected P-wave in X, Y, Z leads
- 15 26. The system according to one of the claims 19 – 25, characterized in that the system is adapted to average the detected P-wave in the predetermined time interval.
- 20 27. The system according to one of the claims 19 – 26, characterized in that the system is adapted to use the first averaged P-wave as a reference P-wave and to compare the upcoming averaged P-waves to it.
- 25 28. The system according to one of the claims 19 – 27, characterized in that the system is adapted to detect at least one loop of the P-wave.
29. The system according to one of the claims 19 – 28, characterized in that, the system is adapted to acquire the ECG-data from a Frank system or a 12-lead ECG-arrangement.
- 30 30. The system according to one of the claims 19 – 29, characterized in that the system is adapted to acquire the ECG-signal from a data storage unit that is independent of this invention and commercially available.

31. The system according to one of the claims 19 – 30, characterized in that the system is adapted to display results of the parameters calculated in trend curve.

5 32. A computer program product, comprising a computer readable storage medium on which is stored a computer program code for a cardiac analysis, which computer program code comprises first computer instructions configured to acquire the ECG-signal, second
10 computer instructions configured to detect at least one wave from the ECG-signal and third computer instructions configured to calculate parameter values of said wave, characterized in that, said wave is P-wave, whereupon the computer program code has instructions for focusing to the dynamic changes of the configuration of said P-wave,
15 wherein said computer program code additionally comprises computer instructions configured to compare substantially every detected P-wave to a reference P-wave in defined time period.

20 33. The computer program product according to claim 32, characterized in that, the cardiac analysis is focused also to dynamic changes of the PQ-segment.

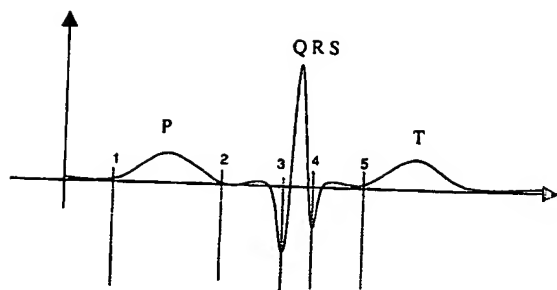


Fig. 1

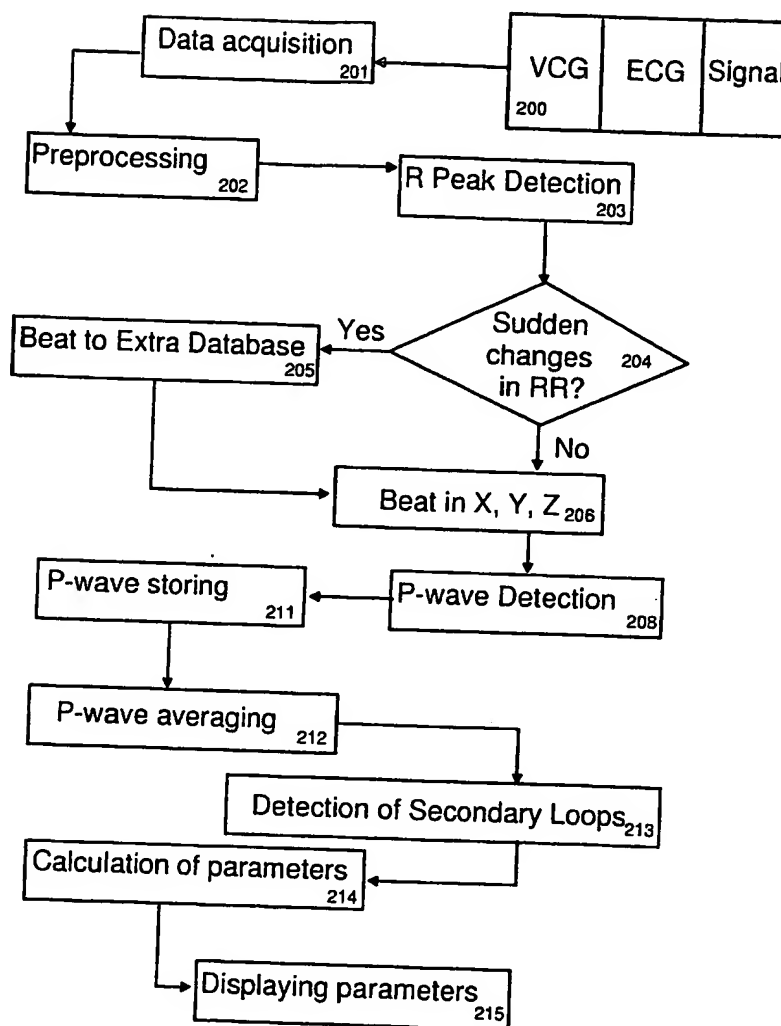


Fig. 2

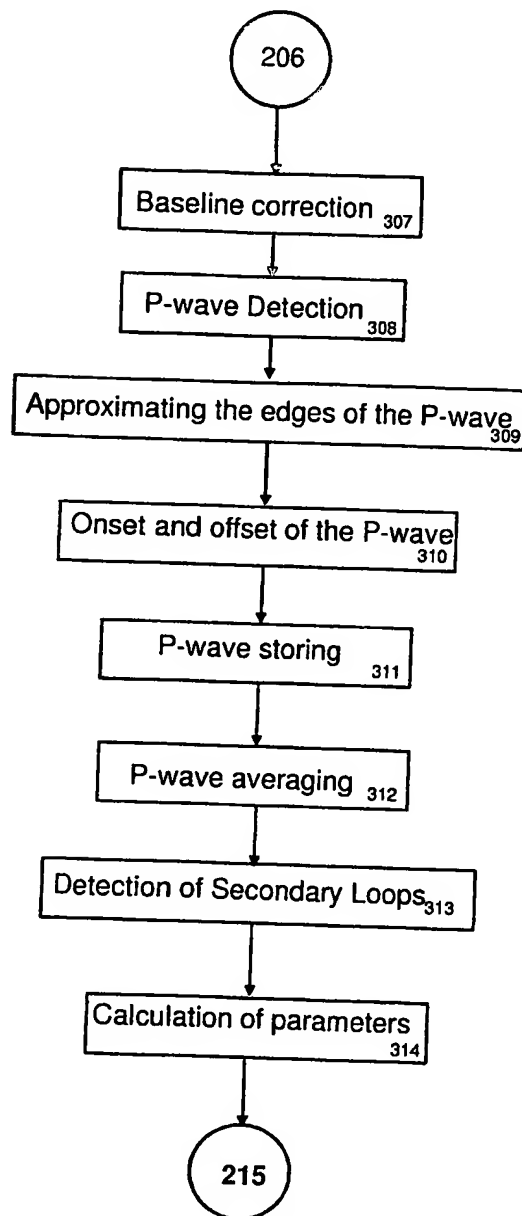


Fig. 3

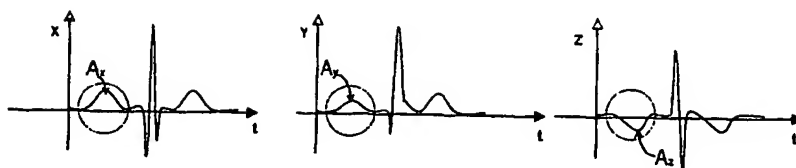


Fig. 4a

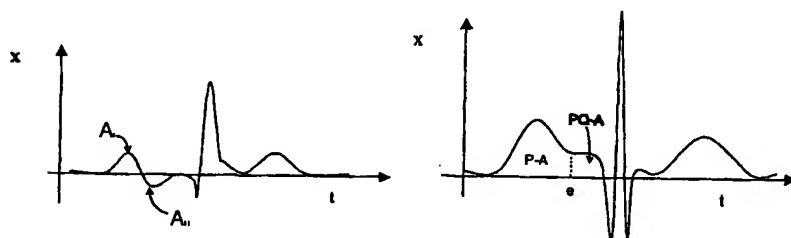


Fig. 4b

Fig. 4c

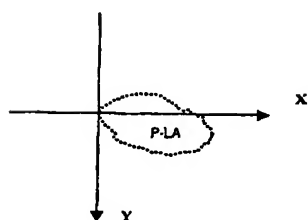


Fig. 5a

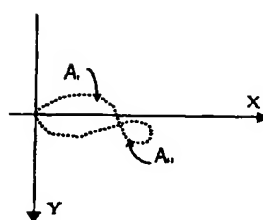


Fig. 5b

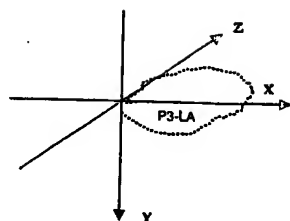


Fig. 6a

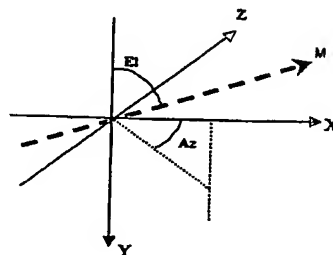


Fig. 6b

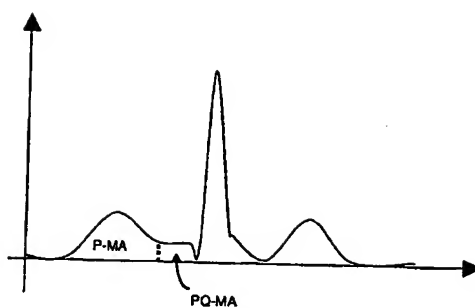


Fig. 7

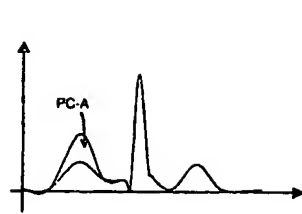


Fig. 8a

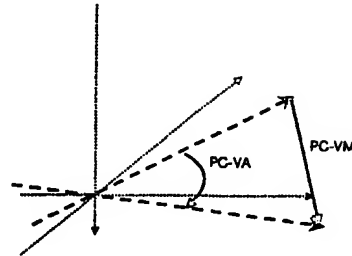


Fig. 8b

INTERNATIONAL SEARCH REPORT

International application No.

PCT/FI 2004/050034

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: A61B 5/0452

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-INTERNAL, WPI DATA

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 0176461 A2 (THE REGENTS OF THE UNIVERSITY OF CALIFORNIA), 18 October 2001 (18.10.2001), page 2, line 7 - line 11; page 7, line 12 - page 8, line 25, figures 1,2 --	1-33
A	US 5755739 A (SUN, W ET AL), 26 May 1998 (26.05.1998) --	1-33
A	US 6516225 A (FLORIO, J J), 4 February 2003 (04.02.2003), abstract -- -----	1-33



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

17 August 2004

Date of mailing of the international search report

20-08-2004

Name and mailing address of the ISA/
Swedish Patent Office
Box 5055, S-102 42 STOCKHOLM
Facsimile No. +46 8 666 02 86

Authorized officer

Patrik Blidefalk /LR
Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.
PCT/FI2004/050034

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: **1 - 18**
because they relate to subject matter not required to be searched by this Authority, namely:
see extra sheet
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/FI2004/050034

Claims 1-18 relates to a method of treatment of the human body by surgery or by therapy/a diagnostic method practised on the human or animal body/Rule 39.1(iv). Nevertheless a search has been executed for these claims. The search has been based on the alleged effects of the product/device.

INTERNATIONAL SEARCH REPORT

Information on patent family members

03/07/2004

International application No.

PCT/FI 2004/050034

WO	0176461	A2	18/10/2001	AU	5386101	A	24/09/2001
				AU	5705201	A	23/10/2001
				US	6556860	B	29/04/2003
				US	6615075	B	02/09/2003
				US	2001056245	A	27/12/2001
				US	2002026220	A	28/02/2002
				WO	0167950	A	20/09/2001
				US	2001056289	A	27/12/2001

US	5755739	A	26/05/1998	NONE			
----	---------	---	------------	------	--	--	--

US	6516225	A	04/02/2003	EP	1123716	A	16/08/2001
				EP	1112757	A	04/07/2001
